Abstract

The present invention relates to hydantoins of formula I,

$$CF_3$$
 $N-R$
 N

in which R is the residue of an am ino carboxylic acid or of an amino carboxylic acid derivative, which is obtained formally by removing an NH₂ group from an amino carboxylic acid or an amino carboxylic acid derivative, to the preparation thereof and to the use thereof as intermediates, in particular for preparing pharmaceutically active ingredients.